



## General

### Guideline Title

Gastrointestinal disorders.

## Bibliographic Source(s)

American Medical Directors Association (AMDA). Gastrointestinal disorders. Columbia (MD): American Medical Directors Association (AMDA); 2006. 28 p. [24 references]

### Guideline Status

This is the current release of the guideline.

This guideline was reaffirmed for currency by the developer in 2011.

# Regulatory Alert

# FDA Warning/Regulatory Alert

Note from the National Guideline Clearinghouse: This guideline references a drug(s) for which important revised regulatory and/or warning information has been released.

March 22, 2016 – Opioid pain medicines
 : The U.S. Food and Drug Administration (FDA) is warning about several safety issues with the entire class of opioid pain medicines. These safety risks are potentially harmful interactions with numerous other medications, problems with the adrenal glands, and decreased sex hormone levels. They are requiring changes to the labels of all opioid drugs to warn about these risks.

# Recommendations

# Major Recommendations

The algorithm Common Gastrointestinal Disorders in the Long-Term Care Setting is to be used in conjunction with the clinical practice guideline. The numbers next to the different components of the algorithm correspond with the steps in the text. Refer to the "Guideline Availability" field for information on obtaining the full text guideline.

# Clinical Algorithm(s)

A clinical algorithm is provided for Common Gastrointestinal Disorders in the Long-Term Care Setting.

# Scope

## Disease/Condition(s)

Gastrointestinal disorders most commonly seen in the long-term care population, including abdominal pain, gastroesophageal reflux disease (GERD), constipation, diarrhea, and gastrointestinal bleeding

Note: Hepatobiliary and pancreatic diseases are beyond the scope of this guideline.

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Diagnosis

Evaluation

Management

Prevention

Risk Assessment

Treatment

## Clinical Specialty

Family Practice

Gastroenterology

Geriatrics

Internal Medicine

Preventive Medicine

### **Intended Users**

Advanced Practice Nurses

Allied Health Personnel

Dietitians

Nurses

Pharmacists

Physician Assistants

Physicians

Social Workers

## Guideline Objective(s)

To improve the quality of care delivered to patients in long-term care settings

To offer care providers and practitioners in long-term care facilities a systematic approach to recognizing, assessing, treating, and monitoring patients with common gastrointestinal disorders

# **Target Population**

Elderly individuals and/or residents of long-term care facilities with common gastrointestinal disorders

### **Interventions and Practices Considered**

#### Recognition/Assessment

Identifying presence of gastrointestinal (GI) disorders by evaluating signs and symptoms and asking questions

Assessing risk factors for GI disorders

Determining the nature and severity of GI disorders using such tools as American Medical Directors Association (AMDA)'s Protocols for

Physician Notification and the PQRST Mnemonic

Identifying the cause of GI disorder (comprehensive history, physical examination, and laboratory tests)

Determining if a referral for a specialty consultation is appropriate

### Management/Treatment/Prevention

Managing the GI disorder and its underlying cause

Identifying and implementing measures to prevent or minimize the risk of GI disorders

#### Monitoring

Monitoring the patient's response to treatment and adjusting interventions as necessary

Monitoring the status and treatment of underlying causes of GI disorders and reviewing relevant medications

Monitoring the facility's management of GI disorders

## Major Outcomes Considered

Signs and symptoms of gastrointestinal (GI) disorders

Risk factors for GI disorders

Use of medications to treat GI disorders

Number of specialist referrals and invasive testing

Morbidity, mortality, and incidence of complications of GI disorders

Incidence of GI disorders

Quality of life

# Methodology

### Methods Used to Collect/Select the Evidence

Searches of Electronic Databases

## Description of Methods Used to Collect/Select the Evidence

2006 Guideline

MEDLINE and PubMed were searched for updated literature related to the subject published between June 2009 and January 2011. This search is done annually and completed by the clinical practice committee vice-chair. If new literature does not change the content or scope of the original guideline, it is deemed to be current. Number of Source Documents Not stated Methods Used to Assess the Quality and Strength of the Evidence Expert Consensus Rating Scheme for the Strength of the Evidence Not applicable Methods Used to Analyze the Evidence Review Description of the Methods Used to Analyze the Evidence Not stated Methods Used to Formulate the Recommendations **Expert Consensus** Description of Methods Used to Formulate the Recommendations This guideline was developed by an interdisciplinary workgroup, using a process that combined evidence and consensus-based approaches. Workgroups include practitioners and others involved in patient care in long-term care facilities. Beginning with a general guideline developed by an agency, association, or organization such as the Agency for Healthcare Research and Quality (AHRQ), pertinent articles and information, and a

draft outline, each group works to make a concise, usable guideline that is tailored to the long-term care setting. Because scientific research in the

long-term care population is limited, many recommendations are based on the expert opinion of practitioners in the field.

# Rating Scheme for the Strength of the Recommendations

Not applicable

Not stated

2011 Review Process

## Cost Analysis

A formal cost analysis was not performed and published cost analyses were not reviewed.

### Method of Guideline Validation

Internal Peer Review

## Description of Method of Guideline Validation

Guideline revisions are completed under the direction of the Clinical Practice Guideline Steering Committee. The committee incorporates information published in peer-reviewed journals after the original guidelines appeared, as well as comments and recommendations not only from experts in the field addressed by the guideline but also from "hands-on" long-term care practitioners and staff.

All American Medical Directors Association (AMDA) clinical practice guidelines undergo external review. The draft guideline is sent to approximately 175+ reviewers. These reviewers include AMDA physician members and independent physicians, specialists, and organizations that are knowledgeable of the guideline topic and the long-term care setting.

# **Evidence Supporting the Recommendations**

## Type of Evidence Supporting the Recommendations

The type of evidence supporting the recommendations is not specifically stated.

The guideline was developed by an interdisciplinary work group using a process that combined evidence- and consensus-based thinking.

# Benefits/Harms of Implementing the Guideline Recommendations

### Potential Benefits

Outcomes that may be expected from implementation of this clinical practice guideline include:

Reduced incidence of some acute gastrointestinal (GI) disorders and greater stability of chronic GI disorders

Appropriate use of medications to treat GI disorders

Appropriate use of acute care facilities to assess and treat GI disorders if indicated

Appropriate use of specialist referrals and invasive testing in the management of GI disorders

Reduced morbidity, mortality, and incidence of complications (e.g., fecal impaction, dehydration) of GI conditions

Improved palliative care outcomes in residents with a poor prognosis

### **Potential Harms**

Adverse Effects of Treatments and Medications

Table 12 in the original guideline document lists adverse effects that should be monitored for.

Both an increased relative risk for community-acquired pneumonia and an increased risk of community-acquired *Clostridium difficile*-associated disease have been reported in patients taking protein pump inhibitors (PPIs). It is uncertain whether the association with pneumonia is caused by the drug or is coincidental to increased use of PPIs in patients with chronic pulmonary conditions who are at increased risk for pneumonia.

Prokinetic agents such as metoclopramide may cause excessive sedation, depression, and tardive dyskinesia.

Any opioid can exacerbate ileus that is causing abdominal pain and should be used with caution.

Both patients and dispensing staff are at some risk of allergy to psyllium.

Because fiber can increase flatulence and fecal incontinence, dosing should be individualized.

Excessive long-term use of stimulant laxatives (e.g., senna, bisacodyl) may be associated with the development of "cathartic colon," that is, a poorly functioning colon caused by the chronic abuse of stimulant laxatives.

The most common side effects of metronidazole are headache, dizziness, GI discomfort, nausea and vomiting, metallic taste, diarrhea,

vaginitis, pelvic discomfort, a disulfiram-like reaction to alcohol, seizures, and peripheral neuropathy. An elevated white blood cell count has also been reported.

Tetracycline may cause a photosensitivity reaction.

Amoxicillin may cause diarrhea or allergy.

## Contraindications

#### Contraindications

Morphine is contraindicated to treat pain caused by cystic duct obstruction or spasm.

Sucralfate is contraindicated if potential exists for significant drug-drug interactions (e.g., phenytoin, thyroid hormone, warfarin)

# Qualifying Statements

## **Qualifying Statements**

This clinical practice guideline is provided for discussion and educational purposes only and should not be used or in any way relied upon without consultation with and supervision of a qualified physician based on the case history and medical condition of a particular patient. The American Medical Directors Association and the American Health Care Association, their heirs, executors, administrators, successors, and assigns hereby disclaim any and all liability for damages of whatever kind resulting from the use, negligent or otherwise, of this clinical practice guideline.

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# Implementation of the Guideline

# Description of Implementation Strategy

The implementation of this clinical practice guideline (CPG) is outlined in four phases. Each phase presents a series of steps, which should be carried out in the process of implementing the practices presented in this guideline. Each phase is summarized below.

#### Recognition

Define the area of improvement and determine if there is a CPG available for the defined area. Then evaluate the pertinence and feasibility of implementing the CPG

#### Assessment

Define the functions necessary for implementation and then educate and train staff. Assess and document performance and outcome indicators and then develop a system to measure outcomes

#### Implementation

Identify and document how each step of the CPG will be carried out and develop an implementation timetable

Identify individual responsible for each step of the CPG

Identify support systems that impact the direct care

Educate and train appropriate individuals in specific CPG implementation and then implement the CPG

#### Monitoring

Evaluate performance based on relevant indicators and identify areas for improvement

Evaluate the predefined performance measures and obtain and provide feedback

# Implementation Tools

Audit Criteria/Indicators

Clinical Algorithm

Tool Kits

For information about availability, see the Availability of Companion Documents and Patient Resources fields below.

# Institute of Medicine (IOM) National Healthcare Quality Report Categories

### IOM Care Need

Getting Better

Staying Healthy

### **IOM Domain**

Effectiveness

# Identifying Information and Availability

## Bibliographic Source(s)

American Medical Directors Association (AMDA). Gastrointestinal disorders. Columbia (MD): American Medical Directors Association (AMDA); 2006. 28 p. [24 references]

## Adaptation

Not applicable: The guideline was not adapted from another source.

### Date Released

2006 (reaffirmed 2011)

## Guideline Developer(s)

American Medical Directors Association - Professional Association

## Guideline Developer Comment

Organizational participants included:

American Association of Homes and Services for the Aging American College of Health Care Administrators American Geriatrics Society

American Health Care Association

American Society of Consultant Pharmacists

National Association of Directors of Nursing Administration in Long-Term Care

National Association of Geriatric Nursing Assistants

National Conference of Gerontological Nurse Practitioners

## Source(s) of Funding

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### Guideline Committee

Steering Committee

## Composition of Group That Authored the Guideline

Committee Members: Marjorie Berleth, MSHA, RNC, FADONA; Lisa Cantrell, RN, C; Charles Cefalu, MD, MS; Sandra Fitzler, RN; Joseph Gruber, RPh, FASCP, CGP; Susan M. Levy, MD, CMD; Harlan Martin, RPh., CCP, FASCP; Evvie F. Munley; Jonathan Musher, MD, CMD; Mary Tellis-Nayak RN, MSN; Barbara Resnick, PhD, CRNP; William Simonson, PharmD., FASCP

### Financial Disclosures/Conflicts of Interest

Not stated

### Guideline Status

This is the current release of the guideline.

This guideline was reaffirmed for currency by the developer in 2011.

# Guideline Availability

Electronic copies: None available

Print copies: Available from the American Medical Directors Association, 10480 Little Patuxent Pkwy, Suite 760, Columbia, MD 21044.

Telephone: (800) 876-2632 or (410) 740-9743; Fax (410) 740-4572. Web site: www.amda.com

# Availability of Companion Documents

The following are available:

Guideline implementation: clinical practice guidelines. Columbia, MD: American Medical Directors Association, 1998, 28 p. We care: implementing clinical practice guidelines tool kit. Columbia, MD: American Medical Directors Association, 2003

Electronic copies: None available

Print copies: Available from the American Medical Directors Association, 10480 Little Patuxent Pkwy, Suite 760, Columbia, MD 21044.

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Additionally, process and quality indicators can be found in Table 16 of the original guideline document.

### Patient Resources

None available

### **NGC Status**

This summary was completed by ECRI on June 23, 2006. This summary was updated by ECRI Institute on April 1, 2009 following the FDA advisory on Reglan (metoclopramide). The currency of the guideline was reaffirmed by the developer in 2011 and this summary was updated by ECRI Institute on May 11, 2012. This summary was updated by ECRI Institute on June 1, 2016 following the U.S. Food and Drug Administration advisory on Opioid pain medicines.

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